

The sources and sinks of CO₂ in caves under mixed woodland and grassland vegetation

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Abstract

We measured concentrations and stable carbon isotope compositions of carbon dioxide in the atmospheres of three caves in central Texas and one cave in southern Arizona in order to identify CO₂ sources and sinks. The vegetation above the caves studied is either savannah (two caves, above which vegetation has been minimally disturbed) or discrete patches of grassland and woodland (two caves, above which vegetation has been highly disturbed). We tested two hypotheses concerning CO₂ in the cave atmospheres: (1) cave ventilation by tropospheric air is the primary sink for CO₂ and (2) CO₂ is primarily derived from the deepest rooting plants growing above the caves. Within caves, we monitored CO₂ at individual locations on monthly and daily time-scales and measured CO₂ along transects with increasing distance from the cave entrances. We also measured CO₂ in the pore spaces of soils under grasses and trees above each of the caves. We calculated $\delta^{13}\text{C}_r$ values of respired CO₂ ($\delta^{13}\text{C}_r$) for all gas samples using measured $\delta^{13}\text{C}$ values and CO₂ concentrations. We then identified the sources of cave CO₂ by comparing cave-air and soil CO₂ $\delta^{13}\text{C}_r$ values. At all locations in each Texas cave, CO₂ concentrations were highest (lowest) and $\delta^{13}\text{C}$ values were lowest (highest) during the summer (winter). Cave-air CO₂ concentrations consistently increased and $\delta^{13}\text{C}$ values consistently decreased with distance from the cave entrances. Similar but smaller magnitude seasonal variations in CO₂ concentrations occurred in the Arizona cave and no seasonal or spatial variation in the $\delta^{13}\text{C}$ of cave-air CO₂ was observed. The mean $\delta^{13}\text{C}_r$ values of CO₂ in soils under grass were 3.5–4.5‰ higher than the $\delta^{13}\text{C}_r$ values of CO₂ in soils under trees. In the caves under savannah, mean $\delta^{13}\text{C}_r$ values of cave-air CO₂ (–24‰ in both caves) were within 1‰ of the mean $\delta^{13}\text{C}_r$ values of CO₂ in soils under trees. In caves covered by large, contiguous areas of grassland, the $\delta^{13}\text{C}_r$ values of cave-air CO₂ were similar to grassland soil values during the summer and were intermediate between grassland and woodland soil values during the winter. The observed spatial and temporal variations in cave-air CO₂ are consistent with density-driven ventilation controlled by seasonal surface temperature changes as the primary sink for CO₂ in the Texas caves. The consistent agreement between soil and cave $\delta^{13}\text{C}_r$ values indicate that the same mixing and diffusion equations that are used to calculate $\delta^{13}\text{C}_r$ values of soil CO₂ also apply to cave-air CO₂. Our results suggest that the majority of CO₂ advects or diffuses into these caves from soils as a gas rather than being transported in aqueous solution. Measured $\delta^{13}\text{C}_r$ values and numerical production-diffusion modeling supports our hypothesis that the majority of gaseous CO₂ in these caves is derived from deeply rooted vegetation. The carbon isotope composition of groundwater and speleothem calcite used for paleoclimate records are therefore likely biased toward deeply rooted plants, even if sparsely present.

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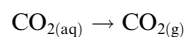
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1. INTRODUCTION

Caves facilitate the study of physical, chemical and biological processes in the deeper portions of the vadose zone. Caves also host speleothems, which preserve geographically widespread, high temporal resolution, and relatively long term (10^3 – 10^4 yrs) Quaternary paleoclimate records that can be dated accurately using U-series disequilibrium methods (see work by L. Edwards, e.g. [Musgrove et al., 2001](#); [Wang et al., 2001](#); [Dykoski et al., 2005](#); and by others, e.g. [Gascoyne, 1992](#); [Richards and Dorale, 2003](#)). We investigated spatial and temporal variations in CO_2 in cave-air to improve understanding of subsurface carbon cycle processes and thereby advance the use of speleothems as paleoclimate indicators.

Previous work has shown that CO_2 concentrations in some caves decrease when ambient atmospheric air temperature drops below cave-air temperature, causing outside air to sink into and displace air from caves (density-driven ventilation, e.g. [Wigley and Brown, 1976](#)). This decrease in cave-air $p\text{CO}_2$ drives degassing of CO_2 dissolved in cave waters and thus the precipitation of calcite according to the following reactions:



Calcite precipitation is suppressed and some speleothems may dissolve when ambient air temperature exceeds cave-air temperature causing ventilation to stagnate and cave-air CO_2 concentrations to increase. Seasonal changes in cave-air CO_2 concentrations are therefore thought to regulate calcite precipitation, and thus speleothem growth rate (e.g. [Spötl et al., 2005](#); [Banner et al., 2007](#); [Baldini et al., 2008](#); [Oster et al., 2012](#)), which is commonly used as a proxy for rainfall. One of our objectives in the present study is to better understand cave CO_2 budgets as a means to improve this proxy with a better understanding of cave CO_2 budgets.

The carbon isotope composition of speleothem calcite is another speleothem-based climate proxy, which may record vegetation changes (e.g. [Dorale et al., 1992, 1998](#); [Hellstrom et al., 1998](#); [Bar-Matthews et al., 1999](#); [Genty et al., 2003](#); [Denniston et al., 2007](#); [Oster et al., 2009](#)) and/or climate-related abiotic changes (e.g. [Dulinski and Rozanski, 1990](#); [Baker et al., 1997](#); [Bar-Matthews et al., 1999](#); [Mickler et al., 2006](#); [Denniston et al., 2007](#); [Oster et al., 2009](#)). $\delta^{13}\text{C}$ values of speleothem calcite, however, are generally considered to be difficult to interpret and are not always reported in isotopic studies of speleothem calcite. A second objective of this study is to advance the interpretation of speleothem calcite $\delta^{13}\text{C}$ values by identifying the sources of carbon in cave-air CO_2 and by extension the source of carbon in epikarst water and speleothem calcite. To this end, we investigate the sources and sinks of cave-air CO_2 by monitoring the concentration and stable carbon isotope composition of cave-air CO_2 and of CO_2 in the pore spaces of soils dominated by grasses and by trees above caves in central Texas and southern Arizona.

2. BACKGROUND

Speleothems archive abundant paleoclimate information. The growth rate, trace element compositions, and stable isotope compositions of calcium carbonate and of fluid inclusions have all been used in numerous studies as paleoclimate proxies ([Fairchild et al., 2006](#) and references cited therein). In particular, speleothem records from central Texas and southern Arizona (regions studied here) show that increased aridity in the southwestern US was associated with transition from the Pleistocene to Holocene warming in the North Atlantic during the last glacial period ([Musgrove et al., 2001](#); [Wagner et al., 2010](#); [Meyer, 2011](#)). In addition, the radiocarbon timescale has been calibrated by comparing U–Th and radiocarbon ages of speleothem calcite ([Hoffman et al., 2010](#); [Southon et al., 2012](#)). Accurate radiocarbon timescale calibrations and proxy-based paleoclimate reconstructions require an understanding of the delivery and removal of CO_2 to and from caves. Specifically, this understanding is key to interpretation of speleothem carbon isotope variations as proxies for temporal shifts in vegetation ([Dorale et al., 1992](#)). Our contribution to this understanding in the present study builds on previous investigations summarized below of carbon cycling through soils, epikarst and caves and the influence that these processes have on speleothem formation and composition.

2.1. Cave CO_2 sinks

Ventilation is perhaps the best-studied process involved in the cycling of carbon through caves and is considered to be the primary mechanism of CO_2 removal from caves (de [Freitas et al., 1982](#); [Troester and White, 1984](#); [Buecher, 1999](#); [Spötl et al., 2005](#); [Baldini et al., 2006, 2008](#); [Banner et al., 2007](#)). While cave-air ventilation may be controlled by many factors, two dominant mechanisms of removal include temperature-driven air flow and wind-driven air flow. The relative importance of these mechanisms varies according to the location and altitude of the entrances and the geometry and volume of the cave ([Buecher, 1999](#); [Fairchild et al., 2006](#); [Batiot-Guilhe et al., 2007](#); [Cowan et al., in press](#)). Although surface temperatures fluctuate seasonally, temperatures within most caves remain nearly constant year-round. Seasonal overturn of cave-air occurs when surface temperatures decrease below cave temperatures, resulting in an unstable density gradient (e.g. Obir Cave, Austria, [Spötl et al., 2005](#); Caves IS and NB of this study, [Banner et al., 2007](#)) Wind-driven ventilation operates according to the Venturi effect: seasonal changes in the direction of wind flowing past the cave entrance, and corresponding changes in wind strength, cause pressure changes below-ground resulting in air flow within the cave ([Kowalczyk and Froelich, 2010](#)). Differences in the gradient and orientation of cave entrances as well as the dominant wind direction influence the cave-troposphere gas exchange. Shorter-term cave-air ventilation has been shown to be correlated to changes in atmospheric pressure, with fluctuations corresponding to thermally driven barometric tides ([Baldini et al., 2006](#); [Cowan, 2010](#)). Similar to the ventilation mech-

anisms described above, the changes in the pressure gradient from the surface to the cave atmosphere drive flow in and out of caves resulting in decreases and increases in cave-air CO₂ concentrations, respectively.

2.2. Cave CO₂ sources

Potential sources of CO₂ in cave-air include: (1) deep geologic sources (magmatic/metamorphic), (2) decomposition of organic matter in caves, (3) local (above cave) soil respiration, (4) animal respiration, and (5) degassing from CO₂-rich groundwater (James, 1977; Troester and White, 1984). Deep sources are not considered in this study because the regions proximal to each cave are not magmatically or tectonically active. Tourism can increase CO₂ concentrations in cave-air by an order of magnitude (Baker and Genty, 1998 and references therein).

In many caves, the dominant sources of CO₂ are atmospheric air and soil respiration (Troester and White, 1984; Ek and Gewalt, 1985; White, 1988; Baldini et al., 2006, 2008). CO₂ from soil can be transported into the caves as a gas or dissolved in seepage waters. Ek and Gewalt (1985) concluded based on spatial variation of CO₂ concentrations that CO₂ produced in soils enters Belgian caves in seepage waters and as a gas through cracks in the epikarst. Spötl et al. (2005) reached the same conclusions for Obir cave, Northern Karawanken Mountains, southern Austria based on stable carbon isotope mass balance calculations. In contrast, stable carbon isotope mass balance suggests CO₂ enters Grotto di Ernesto cave, Asiago–Lavarone karst plateau, northeast Italy primarily as a gas rather than dissolved in seepage water (Frisia et al., 2011). Troester and White (1984) concluded based on similar seasonal variations in CO_{2(g)} and CO_{2(aq)} that CO₂ degassing from an underground stream is the primary source of cave-air CO₂ in Tytoona cave, Pennsylvania, USA.

2.3. Application of carbon isotope ratios to constraining sources and sinks

If soils are the dominant source of CO₂ in most caves, then the theory behind soil CO₂ and its carbon isotope composition should be considered when interpreting measured concentrations and $\delta^{13}\text{C}$ values of cave-air CO₂, and arguably also when interpreting measured $\delta^{13}\text{C}$ values of speleothem calcite. Soil respiration is the sum of CO₂ produced by root/rhizosphere respiration and by the microbial decomposition of organic matter. Respired CO₂ mixes with tropospheric CO₂ belowground in soil pore spaces and accumulates to concentrations that range from hundreds of ppmV to 20%, depending on temperature, water content, porosity, depth and other factors. Accumulation of respired CO₂ in soil pore spaces causes net diffusion of CO₂ along concentration gradients, eventually out of soils into the troposphere. ¹²CO₂ diffuses more rapidly than ¹³CO₂, which in addition to mixing with tropospheric CO₂, results in $\delta^{13}\text{C}$ values of soil CO₂ ($\delta^{13}\text{C}_s$, CO₂ in soil pore spaces) that are at least ~4.4‰ higher than the $\delta^{13}\text{C}$ value of respired CO₂ ($\delta^{13}\text{C}_r$, CO₂ emitted from the soil surface or produced within the soil) (Cerling, 1984; Cerling et al., 1991). At

steady-state, the difference between $\delta^{13}\text{C}_s$ and $\delta^{13}\text{C}_r$ is ~4.4–5.0‰ when soil CO₂ concentrations are above ~10,000 ppmV. The difference between $\delta^{13}\text{C}_s$ and $\delta^{13}\text{C}_r$ increases as soil CO₂ concentrations decrease such that $\delta^{13}\text{C}_s - \delta^{13}\text{C}_r$ can be as large as 11‰ at 1000 ppmV. This effect complicates interpretations of the source of respired CO₂ based on $\delta^{13}\text{C}_s$ values alone. To circumvent this complexity, we solved the following equation derived by Davidson (1995) for $\delta^{13}\text{C}_r$:

$$\delta_s = 1.0044\delta_r + \frac{C_a}{C_s}(\delta_a - 1.0044\delta_r - 4.4) + 4.4 \quad (1)$$

where δ is the $\delta^{13}\text{C}$ value, C is the ¹²CO₂ concentration which can be approximated by the total CO₂ concentration, and the subscripts s, a and r represent soil, atmospheric and respired CO₂, respectively. We use the rearranged form of Eq. (1) to assess the relative influence of tree- and grass-dominated soils on cave-air CO₂.

3. STUDY SITES

Three caves located on the Edwards Plateau in central Texas and one cave located on the southeastern flank of the Santa Rita mountains in southern Arizona were studied: Inner Space Cavern (IS), Natural Bridge Caverns (NB) and Whirlpool Cave (WP) in Texas, and Cave of the Bells (COB) in Arizona. The land surfaces above some of the caves in this study have been affected by the construction of roads and buildings. The area on the surface around IS is the most urbanized of all locations; NB and WP are located in less disturbed settings; COB is remote and undisturbed. Satellite images of the distribution of vegetation above each cave are shown in Fig. 1. The passages extend down from the entrances of all the caves studied.

The Texas caves are situated within approximately 150 km of each other and have similar surface climate. Average surface air temperatures in central Texas are 36 °C in the summer (JJA) and 4 °C in the winter (DJF). The region generally experiences dry summers, with peak rainfall in the spring and, depending on tropical storm and hurricane tracks, in the fall as well. Soils above the Texas caves studied here are Lithic Haplustolls or Lithic Argiustolls and are thin (<30 cm) and rocky with fragments from the underlying bedrock (Godfrey et al., 1973; Cooke et al., 2007). Vegetation above NB is juniper and oak savannah whereas patches of grassland and woodland occur above the other Texas caves. Dominant tree species above the Texas caves are *Juniperus ashei* (ashe juniper) and *Quercus virginiana* (Texas live oak). Grasslands are dominated by *Bothriochloa ischaemum* (King Ranch blue-stem) and *Carex planostachys* (Cedar Sedge) but also contain C₃ plants. Small percentages of the surfaces above each cave are covered by Texas Prickly Pear cactus (*Opuntia engelmannii*) a CAM plant. A study of roots exposed in caves on the Edwards Plateau (including NB studied here) demonstrated that *J. ashei* roots extend at least 8 m into bedrock and that *Q. virginiana* var. *fusiformis* roots extend to ~20 m (Jackson et al., 1999). No roots were found in caves below 25 m depth (Jackson et al., 1999). The majority of grassland root biomass likely occurs within the top

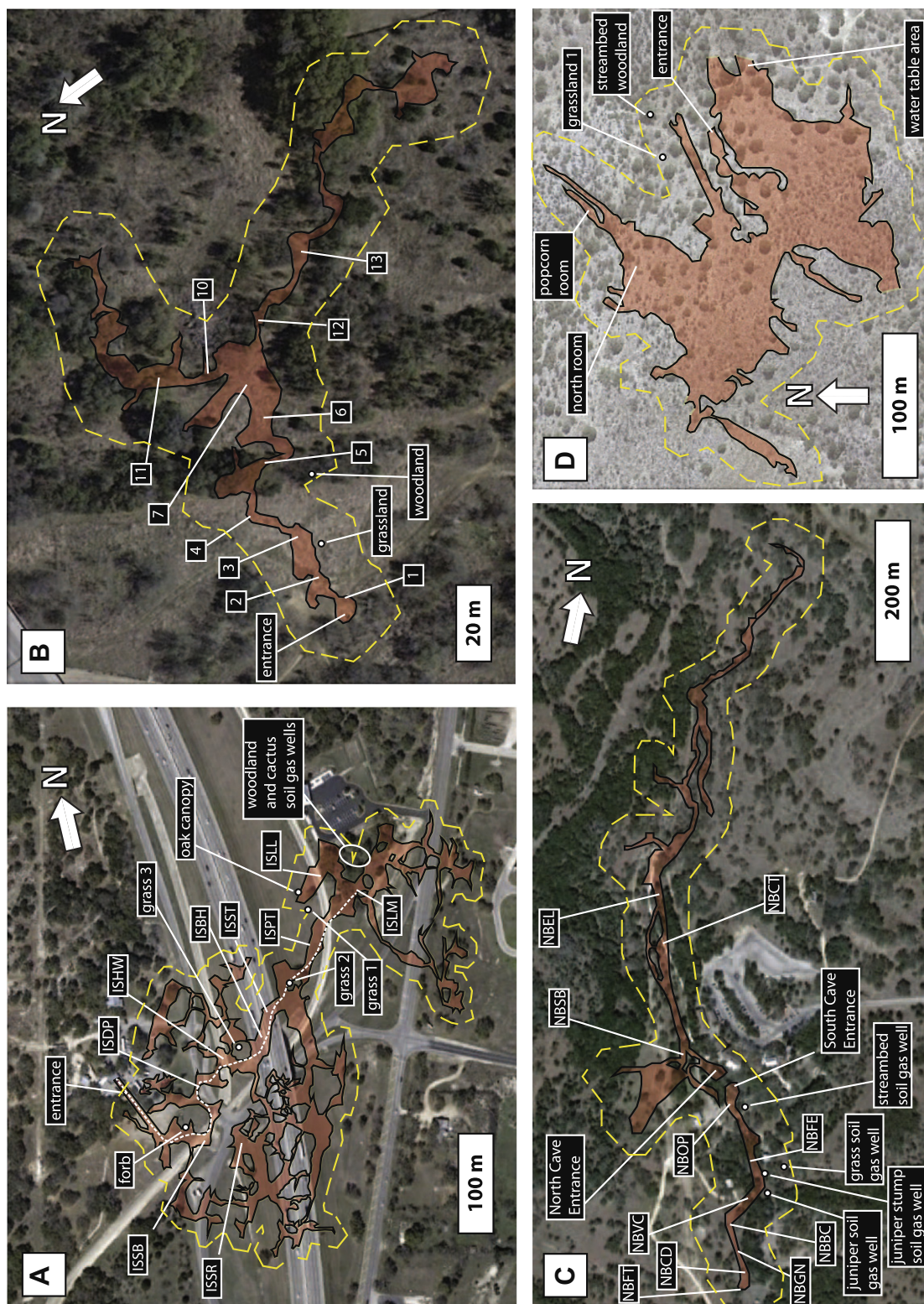


Fig. 1. Aerial view maps of the caves studied. (A) Inner Space Caverns (IS), (B) Whirlpool Cave (WP), (C) Natural Bridge Caverns (NB) and (D) Cave of the Bells (COB). The locations of the caves are shown in orange. Gas sample collection sites in the caves and in the soils above the caves are labeled; soil gas sampling locations are indicated by circles. The dashed yellow lines outline the area above each cave that is within one mean cave depth of the walls of each cave. The areas within the yellow dashed lines were used to calculate the fraction of the surface above the caves covered by tree canopies. IS and COB are covered by 15% tree canopy, WP by 50% tree canopy and NB by 60% tree canopy.

Table 1
Comparison of caves.

Cave name*	Location	Vegetation	Disturbance**	Radius of mean cave			
				Tourist cave	Entrance (m)	Depth (m)	Volume (m ³)
IS	Central TX	Patches of woodland and grassland	High	Yes	3	12	75,000
NB	Central TX	Oak and juniper savannah	Low	Yes	2	40	400,000
COB	Southern AZ	Oak and juniper savannah	Low	No	0.25	20	>40,000
WP	Central TX	Patches of woodland and grassland	High	No	1	10	22,000

* IS = Inner Space Cavern, NB = Natural Bridge Cavern, COB = Cave of the Bells, WP = Whirlpool.

** Degree of anthropogenic disturbance of vegetation.

50 cm; in nearby sandy loams, 80% of the herbaceous root biomass occurred in the top 60 cm of soil (Brown and Archer, 1990). Cactus roots typically do not extend below 30 cm with most roots in top 10 cm (Nobel, 1988).

Southern Arizona has a monsoon climate. Mean annual temperature at Cave of the Bells is about 14 °C with maximum air temperatures of 23 °C and minimum of 5 °C. COB is located beneath a hill flanked by seasonal streams. Soils above COB are shallow and gravelly Entisols and calcic Aridisols. The vegetation above the cave is characterized as juniper – oak savannah on the northeast-facing slope and as desert scrub on the southwest facing slope. *Juniperus deppeana* (alligator juniper) and *Quercus arizonica* (Arizona white oak) are the dominant tree species in the savannah, which also includes *Rhus choriophylla*, *Mimosa biuncifera*, *Bouteloua curtipendula* and *Bothriochloa barbinodis*. The dominant species in the desert scrub include *Viguiera sp.*, *Opuntia discata*, *Cowania mexicana*, *B. curtipendula*, and *Macrosiphonia brachysiphon*. There is a higher density of trees in and along the seasonal streams than on the hillslopes.

Characteristics of the caves studied are summarized in Table 1. Caves IS and NB are the two largest caves in this study. In both caves, public tours operate year-round and the natural entrances to both caves have been altered (Elliot and Veni, 1994; Banner et al., 2007). The caves contain ventilation shafts that are occasionally used in the summer months to decrease the build-up of CO₂ in order to make the environment comfortable for tourism (Cowan, 2010). IS has a total surveyed length of 4851 m, a maximum depth of 24 m, and a total volume of approximately 75,000 m³ (Elliot and Veni, 1994; Cowan, 2010; Fig. 1A). The profile of the cave grades downward from the entrance; the vertical relief is 17 m from the entrance to the rear of the commercial path (Elliot and Veni, 1994). The cave consists of passages roughly 2 m across and numerous chambers greater than 2 m across. The transect sampled in this study traverses one of the larger of these chambers, where sites Borehole (ISBH) and Flowing Stone of Time (ISST) are located. No natural entrance to the cave exists. A large, gradually sloping artificial entrance to the cave faces the NW and is approximately 3 m in radius. NB has a length of 2621 m and a maximum depth of 76 m (Elliot and Veni, 1994; Fig. 1C). The vertical relief of the cave is much larger than that of IS (~50 m). The cave is composed of two caverns with separate entrances, Natural Bridge South (NBS) and Natural Bridge North (NBN). The approximate volume

of NBS is 150,000 m³ and of NBN is 250,000 m³ (Cowan, 2010). Entrances to NBS and NBN are sealed by glass doors and face the WNW and SSE respectively. Vertical ventilation shafts to the surface are located in both caverns.

WP is much smaller in volume than the commercial caves. Passages within this cave are narrow and contain a few small chambers. Public access to WP is restricted. WP has a maximum depth of 13 m and a total volume of approximately 22,000 m³ (Cowan, 2010; Fig. 1B). The nearly vertical entrance to the cave is covered by a gate that does not impede air flow. The profile of the cave beyond the initial drop at the entrance slopes gradually downward towards the east. The cave is Y-shaped with the majority of sampling sites for this study located within the first 1/3 of the cave (Fig. 1B). Passages are approximately 1 m in height and 4 m in width (Elliot and Veni, 1994). Active vadosic drips were not observed in WP during the course of this study.

COB has the most complicated geometry of the caves studied. It has multiple levels and passages that lead up and down repeatedly, which likely limits density-driven ventilation. At least one passage in COB extends down to the water table. The entrance to COB is on a northeast facing hillslope and the passage leading into the cave constricts to an opening approximately 0.5 × 0.5 meters in cross section and approximately 1 m long, which might also restrict ventilation.

4. METHODS

Air sampling sites in the four caves were chosen along spatial transects with increasing distance into the caves. Cave-air samples were obtained on seasonal intervals, except at IS and COB where samples were collected on 4–12 week intervals. In addition to the monthly collection trips, cave-air samples were retrieved from IS every 2-h over a 24-h period between July 14th and July 15th, 2010. We collected samples in duplicate at each site, and the CO₂ concentration and temperature were measured using a portable Telaire 7001 CO₂ meter, with an uncertainty of approximately 5%.

Cave-air and soil-gas samples were collected using plastic disposable syringes and immediately injected into septum-capped, He-flushed 12 mL glass vials (Labco exetainers®) for transport to the laboratory. Cave-air was collected at least 5 m further into the cave than the rest of the group on each cave sampling trip. In addition, in order

to minimize any further anthropogenic effects on the cave-air samples, the person doing the sampling held their breath while taking the air sample. Air samples were also collected at the surface (tropospheric air) using this same procedure during each cave sampling trip. Human breath was sampled by blowing for five seconds into a needle penetrating the septum of a He-flushed exetainer[®]. A second needle was used as a vent to allow breath to displace helium in the vial.

Soil-gas wells of the design described in Breecker and Sharp (2008) were installed above IS, NB, WP and COB. The soil-gas wells were made of stainless steel tubing 40–50 cm in length and 5 mm in diameter. The bottom of the tubes were crimped shut and a slit, approximately 2 mm in width, was filed into the wall of the tubing near the crimp. Wells were inserted vertically into the soil above the caves using a battery-powered drill. A Swagelok 1/4" fitting and rubber septum were placed on the top end of the well to prevent any direct exchange of gas in the well with tropospheric air. Wells were installed in grasslands and woodlands (or under tree canopy in savannah) at depths ranging from 5 cm and 55 cm. Samples were collected by syringe through the rubber septum. A 3 mL (1 mL at COB) syringe was first flushed with gas from the well before samples were collected. A sample was then collected by extracting 3 mL of soil-gas from each well (0.2–2.0 mL for soil gas wells above COB) and injecting it into a He-flushed, septum-capped vial. Duplicate samples were collected from each soil-gas well to evaluate precision. Recoil of the syringe plunger indicated sub-atmospheric pressure in the well and required longer syringe collection times. This procedure ensured that tropospheric air did not contaminate the soil gas samples upon transfer from well to vial.

The soil and cave gas samples were analyzed using continuous flow gas chromatography/isotope ratio mass spectrometry (GC/IRMS). Within a week of collection an aliquot of each sample was transferred to a newly flushed exetainer[®] vial (no transfer was made for samples collected from COB soil gas wells, instead the entire sample was analyzed). Sample air was flushed from the vials in a He stream and directed through a liquid nitrogen cooled trap using a Gasbench II. The CO₂ in each aliquot was cryofocused and analyzed as a single pulse in continuous flow mode on a Delta Plus mass spectrometer. Carbon isotope compositions are expressed in the standard delta notation relative to VPDB. A CO₂-in-air standard calibrated to the VPDB scale in the Stable Isotope Laboratory at CU-INSTAAR in Boulder, Colorado (for calibration methods see Trolier et al., 1996) was used to calibrate an internal laboratory CO₂-in-air standard, which was analyzed alongside samples during each run.

Air samples for radiocarbon measurement were collected in evacuated, 1 L glass flasks. Air was drawn into the flasks from the soil (by connection to a manifold previously flushed with soil gas) or cave atmosphere through a quartz wool filter. The soil gas sample was collected in February 2009 from 55 cm depth in one of the grasslands above COB (grassland 1, Fig. 1). Cave-air was collected in November 2008 in the North Room and in February 2009 just above the water table (Fig. 1). CO₂ was separated

from air cryogenically, passed through silver and copper wool to remove halogens and graphitized for analysis. The fraction modern carbon (fmc), as defined by Stuiver and Polach (1977), in each sample was determined in the Accelerator Mass Spectrometry laboratory at the University of Arizona.

5. RESULTS

The concentrations and $\delta^{13}\text{C}$ values of soil CO₂ and cave-air CO₂ measured in this study are presented in Tables 2–8. The $\delta^{13}\text{C}$ values of CO₂ in IS ranged from -17.7‰ to -8.6‰ and CO₂ concentrations ranged from 340 ppm to 8830 ppm. Spatial transects show a general trend toward decreasing $\delta^{13}\text{C}$ values with distance from the cave entrance (Fig. 2). $\delta^{13}\text{C}$ values of CO₂ from NBS range from -19.1‰ to -15.7‰ and CO₂ concentrations range from 750 to 13,700 ppm. $\delta^{13}\text{C}$ values of CO₂ from NBN range from -18.6‰ to -15.1‰ and CO₂ concentrations range from 750 to 3500 ppm. Whereas the maximum CO₂ concentrations at NBN were significantly lower than those of NBS, the range in the $\delta^{13}\text{C}$ values was comparable. $\delta^{13}\text{C}$ values of CO₂ from WP range from -21.0‰ to -14.9‰ and CO₂ concentrations range from 1400 to 17,800 ppm. $\delta^{13}\text{C}$ values of cave-air CO₂ within WP decrease with distance from the entrance. The $\delta^{13}\text{C}$ values of CO₂ in COB air vary by $\pm 0.4\text{‰}$ (1σ , $n = 21$, mean value = -19.2‰) with no spatial pattern detected. CO₂ concentrations in COB vary from ~ 8000 ppmV in the winter to $\sim 12,000$ ppmV in the summer.

The most complete time series of cave-air CO₂ measurements are from IS and COB. In IS, the lowest CO₂ concentrations and the highest $\delta^{13}\text{C}$ values occurred during the months of December, January and February of each year and the highest CO₂ concentrations and lowest $\delta^{13}\text{C}$ values were measured in June, July, and August (Table 2, Fig. 2). Similar seasonal changes were measured in Obir caves by Spötl et al. (2005). At IS, the magnitude of seasonal variation in CO₂ concentration and $\delta^{13}\text{C}$ values is largest nearest the entrance and decreases with distance into the cave. As a result the decrease of $\delta^{13}\text{C}$ values with distance from the cave entrance is larger during the winter than during the summer (Fig. 2). In COB, no seasonal pattern in $\delta^{13}\text{C}$ values occurs and variations in the $\delta^{13}\text{C}$ values and concentrations of CO₂ are smaller than they are in IS. The fmc in COB cave-air CO₂ was 0.94753 ± 0.00283 and 0.94744 ± 0.00014 in November 2008 and February 2009, respectively, which yielded a conventional radiocarbon age of 430 years BP. The fmc in the grassland soil at COB was 1.08365 ± 0.00016 in February 2009, which is indistinguishable from the present-day tropospheric CO₂ values.

Calculated cave-air CO₂ $\delta^{13}\text{C}_r$ values are listed in Tables 2–8 and are plotted in Fig. 3. Mean $\delta^{13}\text{C}_r$ values of cave-air CO₂ in IS, NB, WP and COB are similar (-24.2‰ , -23.8‰ , -23.1‰ and -23.9‰ , respectively). Cave-air CO₂ $\delta^{13}\text{C}_r$ values vary seasonally in IS and WP; the lowest values occur in the winter or spring and the highest values in July or August. Cave-air CO₂ $\delta^{13}\text{C}_r$ values in NB and COB are less variable. When CO₂ concentrations in IS

Table 2
CO₂ concentrations and $\delta^{13}\text{C}$ values in IS cave-air.

Date	Outside air		ISSB		ISDP		ISHW		ISBH		ISST		ISLM		ISSR		ISLL									
	pCO ₂	δ ¹³ C	pCO ₂	δ ¹³ C	pCO ₂	δ ¹³ C	pCO ₂	δ ¹³ C	pCO ₂	δ ¹³ C	pCO ₂	δ ¹³ C	pCO ₂	δ ¹³ C	pCO ₂	δ ¹³ C	pCO ₂	δ ¹³ C								
7/3/2008	360	-9.2	781	-14.7	-23.7	1455	-13.0	-18.6	1260	-13.8	-20.0	1493	-13.8	-19.6	1654	-13.5	-19.0	2483	-13.5	-18.5	2846	-17.0	-22.4			
7/19/2008	392	-9.3	2878	-15.3	-20.6	2819	-15.3	-20.6	2968	-15.4	-20.6	3154	-15.3	-20.5	3143	-15.3	-20.5	4370	-16.3	-21.3	4478	-16.3	-21.3			
9/23/2008	331	-10.0	1131	-14.7	-21.0	1680	-16.1	-21.9	1311	-15.8	-22.1	1708	-16.2	-22.0	1689	-16.1	-21.9	5307	-18.0	-22.8	4743	-17.6	-22.5			
10/23/2008	300	-8.6	348	-10.5	-26.7	385	-12.2	-29.2	597	-13.1	-21.9	714	-12.7	-20.0	717	-14.2	-22.5	4293	-17.7	-22.7	1467	-16.5	-22.8			
12/6/2008	342	-9.9	339	-10.3		361	-11.2	-38.8	380	-11.2	-27.2	376	-12.5	-42.9	376	-12.1	-38.5	386	-12.9	-40.4	517	-14.3	-27.2	639	-15.6	-26.4
2/15/2009	320	-8.7	362	-11.0	-32.8	411	-14.7	-40.0	398	-12.3	-31.3	382	-11.7	-31.4	343	-13.1	-78.4	446	-12.1	-25.0	485	-13.8	-28.0	670	-15.3	-25.6
5/17/2009	244	-8.4				342	-9.2	-15.5	382	-11.3	-20.7	416	-11.8	-20.9	450	-10.0	-16.2	762	-12.8	-19.2	692	-13.0	-19.8	1080	-14.3	-20.3
8/10/2009	300	-7.9	2535	-15.6	-20.9	2974	-14.1	-19.1	3143	-15.3	-20.4	3840	-15.9	-20.9	4083	-15.9	-20.8	4862	-16.2	-21.1	5742	-15.8	-20.5	4575	-16.4	-21.3
10/4/2009	590	-10.2	1940	-15.6	-22.3	1910	-15.6	-22.3	2030	-15.6	-22.1	2330	-15.9	-22.1	2470	-15.9	-22.0	8830	-17.4	-22.2	4700	-16.3	-21.5	4540	-16.6	-21.9
6/29/2010	490	-9.4				2238	-16.6	-23.0							1652	-16.0	-23.1	3004	-17.0	-22.8	3048	-16.2	-21.8			
7/14/2010	543	-10.0				3319	-16.9	-22.6	3459	-16.7	-22.2				3429	-16.3	-21.8	4779	-17.6	-22.9						

are below 1500 ppmV CO₂, $\delta^{13}\text{C}_r$ values decrease with decreasing cave-air CO₂ concentration (Fig. 3D). During the 24 h study in IS, cave-air CO₂ concentrations reached a minimum at ISPT of 3500 ppmV at 07:00 (CST) and a maximum of 4400 ppm at 13:00 (CST) whereas $\delta^{13}\text{C}_r$ reached a maximum of -21.8‰ at 07:00 (CST) and a minimum of -22.8‰ at 13:00 (CST) (Fig. 4). Along transects into the cave, 1‰ changes in $\delta^{13}\text{C}_r$ values occur over <75 m and the position of these transitions shift in space diurnally (Fig. 4).

Soil CO₂ collected from gas wells above IS, NB, and WP on separate collection trips had $\delta^{13}\text{C}$ values ranging from -25.3‰ to -17.4‰ (Tables 3, 5 and 7; Fig. 3). $\delta^{13}\text{C}_r$ values for woodland and cactus soils above IS are higher in the summer (June, July, August, September) than in the spring (March) and fall (October) by up to 3.5‰. $\delta^{13}\text{C}_r$ values for grassland soils above COB vary from -22‰ during the winter to -19‰ in May and June. $\delta^{13}\text{C}_r$ values in the streambed woodland soil above COB are relatively constant ($-23.9 \pm 0.6\text{‰}$, 1σ , $n = 12$). Mean $\delta^{13}\text{C}_r$ values of soil CO₂ under juniper trees are similar above NB, WP, and COB (-24.1‰ , -23.5‰ and -23.9‰ , respectively). Mean grassland soil $\delta^{13}\text{C}_r$ values are similar at IS, NB, WP, and COB (-20.3‰ , -19.0‰ , -19.7‰ and -20.5‰). There is a close agreement between cave-air CO₂ $\delta^{13}\text{C}_r$ values and tree-dominated soil $\delta^{13}\text{C}_r$ values at NB and COB. Cave-air CO₂ $\delta^{13}\text{C}_r$ values in IS and WP are similar to grassland $\delta^{13}\text{C}_r$ values during the summer and are intermediate between grassland and woodland values during the winter (Fig. 3). The mean $\delta^{13}\text{C}$ value of CO₂ in human breath was -21.5‰ (duplicate samples from five individuals, $1\sigma = 0.6$).

6. DISCUSSION

We apply our results to understanding the processes involved in carbon transport through karst and outline implications for interpretation of speleothem carbon isotope proxies. Cave-air CO₂ measured in this study is a mixture of two endmembers (as inferred by Banner et al., 2007): tropospheric air (~ 390 ppmV, -8.5‰ $\delta^{13}\text{C}$, VPDB) and an endmember with higher CO₂ and lower $\delta^{13}\text{C}$ values. The identification of tropospheric CO₂ as one of the two endmembers is supported by spatial trends of increasing $\delta^{13}\text{C}$ values and decreasing CO₂ concentrations toward the cave entrances and the near tropospheric values near the entrance of IS. Spötl et al. (2005) also interpreted cave-air CO₂ as a two-component mixture (tropospheric CO₂ and a 'light' endmember). The mixing relationships discussed below support density-driven ventilation as the primary sink for cave-air CO₂ and deeply rooted trees as the primary source of carbon in cave-air CO₂.

6.1. Evidence for ventilation

The spatial and temporal correlation between $\delta^{13}\text{C}$ values and concentrations of cave-air CO₂ are consistent with increased mixing ratios of atmospheric CO₂ in these caves during winter (compared with summer) and closer to the

Table 3
CO₂ concentrations and $\delta^{13}\text{C}$ values in IS soils.

Date	Outside air		Soil cactus 1 (28 cm)			Soil cactus 2 (11–26 cm)			Soil oak 1 (25 cm)			Soil oak 2 (7–22 cm)			Soil west (26 cm)		
	$p\text{CO}_2$	$\delta^{13}\text{C}$	$p\text{CO}_2$	$\delta^{13}\text{C}$	$\delta^{13}\text{C}_r$	$p\text{CO}_2$	$\delta^{13}\text{C}$	$\delta^{13}\text{C}_r$	$p\text{CO}_2$	$\delta^{13}\text{C}$	$\delta^{13}\text{C}_r$	$p\text{CO}_2$	$\delta^{13}\text{C}$	$\delta^{13}\text{C}_r$	$p\text{CO}_2$	$\delta^{13}\text{C}$	$\delta^{13}\text{C}_r$
3/27/2010	540	−9.8	9783	−20.4	−25.3				8414	−22	−27.1				10,290	−22	−27.0
6/29/2010	490	−9.4	6830	−18.1	−23.1				3178	−18.5	−24.5				23,216	−20	−24.0
6/27/2011			5375	−15.1	−19.8	13,720	−17.4	−21.9	17,554	−19.4	−23.9				22,527	−25	−29.8
8/16/2011			571	−11.3	−19.5	2875	−17	−22.4	741	−13.2	−21.5	1548	−17.4	−24.3	3772	−18	−22.9
9/9/2011			614	−12.8	−22.5	1398	−16.7	−23.7	517	−12.8	−26.0	987	−17.6	−26.9	1792	−19	−25.4
10/9/2011			2020	−17.7	−24.0	1341	−16.8	−24.2	2144	−20.8	−27.6	998	−17.9	−27.6	2249	−19	−25.3
Date	Outside air		Grassland 1 (10–25 cm)			Grassland 2 (9–24 cm)			Grassland 3 (9–24 cm)			Oak canopy savannah (5–20 cm)			Forb (4–19 cm)		
	$p\text{CO}_2$	$\delta^{13}\text{C}$	$p\text{CO}_2$	$\delta^{13}\text{C}$	$\delta^{13}\text{C}_r$	$p\text{CO}_2$	$\delta^{13}\text{C}$	$\delta^{13}\text{C}_r$	$p\text{CO}_2$	$\delta^{13}\text{C}$	$\delta^{13}\text{C}_r$	$p\text{CO}_2$	$\delta^{13}\text{C}$	$\delta^{13}\text{C}_r$	$p\text{CO}_2$	$\delta^{13}\text{C}$	$\delta^{13}\text{C}_r$
12/20/2011	409	−8.6	1498	−15.0	−21.7	1235	−13.3	−20.0	1132	−12.7	−19.3	3836	−20.5	−26.2	1212	−18	−26.6

Table 4
CO₂ concentrations and $\delta^{13}\text{C}$ values in NB cave-air.

Date	Natural Bridge South														
	Outside air		Step 92			NBOP			NBEP			NBVC			
	$p\text{CO}_2$	$\delta^{13}\text{C}$	$p\text{CO}_2$	$\delta^{13}\text{C}$	$\delta^{13}\text{C}_r$	$p\text{CO}_2$	$\delta^{13}\text{C}$	$\delta^{13}\text{C}_r$	$p\text{CO}_2$	$\delta^{13}\text{C}$	$\delta^{13}\text{C}_r$	$p\text{CO}_2$	$\delta^{13}\text{C}$	$\delta^{13}\text{C}_r$	
6/11/2008	380	−8.5	1752	−18.7	−25.8	1684	−18.4	−25.6	5197	−18.6	−23.7	4701	−17.3	−22.4	
3/8/2009	268	−8.6	870	−17.5	−25.7	762	−15.7	−23.8				2252	−18.5	−24.1	
6/17/2010															
Date	Natural Bridge South														
	NBFE			NBBC			NBGN			NBCD			NBFT		
	$p\text{CO}_2$	$\delta^{13}\text{C}$	$\delta^{13}\text{C}_r$	$p\text{CO}_2$	$\delta^{13}\text{C}$	$\delta^{13}\text{C}_r$	$p\text{CO}_2$	$\delta^{13}\text{C}$	$\delta^{13}\text{C}_r$	$p\text{CO}_2$	$\delta^{13}\text{C}$	$\delta^{13}\text{C}_r$	$p\text{CO}_2$	$\delta^{13}\text{C}$	$\delta^{13}\text{C}_r$
6/11/2008	13,700	−18.4	−23.0	4748	−17.2	−22.3	4790	−19.1	−24.3	6651	−15.8	−20.6	5927	−18.6	−23.6
3/8/2009	1916	−18.1	−23.9	2374	−18.7	−24.3	2372	−18.5	−24.1	2568	−18.7	−24.2	3171	−18.9	−24.1
6/17/2010	4174	−17.3	−22.5							4609	−18.4	−23.6	4565	−18.5	−23.7
Date	Natural Bridge North														
	NBSB			NBCT			NBEL								
	$p\text{CO}_2$	$\delta^{13}\text{C}$	$\delta^{13}\text{C}_r$	$p\text{CO}_2$	$\delta^{13}\text{C}$	$\delta^{13}\text{C}_r$	$p\text{CO}_2$	$\delta^{13}\text{C}$	$\delta^{13}\text{C}_r$	$p\text{CO}_2$	$\delta^{13}\text{C}$	$\delta^{13}\text{C}_r$	$p\text{CO}_2$	$\delta^{13}\text{C}$	$\delta^{13}\text{C}_r$
6/11/2008	3398	−18.5	−24.1				3079	−17.7	−23.3		3213		−18.6		−24.2
3/8/2009	837	−15.4	−22.9				744	−15.1	−23.1		837		−16.6		−24.7
6/17/2010							3560	−17.6	−23.0						

entrances (compared with deeper regions of the caves). These observations confirm inferences made from CO₂ concentration time series that cave atmospheres are ventilated by tropospheric air during the winter (Banner et al., 2007; Cowan et al., in press). The magnitude of the effect of ventilation on cave-air CO₂ varies among the caves studied. Ventilation has a minor effect in COB compared with the other caves; the mixing ratio of tropospheric CO₂ in this cave is too small to have a measurable effect on cave-air CO₂ $\delta^{13}\text{C}$ values, even during the winter. The small area of the entrance and the undulating passages in COB likely limit density-driven ventilation. Ventilation has a larger effect in IS than in any of the other caves studied. The deep ventilation in IS may result from the large cross sectional area of the cave's entrance or possibly from wind blowing into the cave. Northerly winds accompany cold fronts

moving into central Texas during the winter (NOAA) and IS is the only cave studied with a north-facing entrance. Given the resolution of the data from IS, however, it is unclear whether or not northerly winds enhance ventilation.

6.2. The source of carbon in cave-air CO₂

We determine the area above each cave (within the dashed lines in Fig. 1, which correspond to the projected area of each cave expanded in every direction by a distance equivalent to the depth of that cave) covered by grassland and by tree canopies using Adobe Illustrator to calculate the area of irregularly shaped polygons. Tree canopies cover 15% of the surface above IS and COB, 50% of the surface above WP and 60% of the surface above NB. The majority of the remainder of the surface above each cave is covered

Table 5
CO₂ concentrations and $\delta^{13}\text{C}$ values in NB soils.

Date	Juniper														
	10 cm			15 cm			21 cm			24 cm			34 cm		
	$p\text{CO}_2$	$\delta^{13}\text{C}$	$\delta^{13}\text{C}_r$	$p\text{CO}_2$	$\delta^{13}\text{C}$	$\delta^{13}\text{C}_r$	$p\text{CO}_2$	$\delta^{13}\text{C}$	$\delta^{13}\text{C}_r$	$p\text{CO}_2$	$\delta^{13}\text{C}$	$\delta^{13}\text{C}_r$	$p\text{CO}_2$	$\delta^{13}\text{C}$	$\delta^{13}\text{C}_r$
6/17/2010	3102	−18.5	−24.2	5578	−19.4	−24.5	8215	−20	−24.8	11,102	−19.9	−24.6	18,896	−20.1	−24.6
6/27/2011	2565	−18.1	−24.1	3037	−18.6	−24.3	6068	−19.3	−24.3	15,547	−20	−24.6	6284	−19.5	−24.5
8/19/2011	1104	−15.6	−23.6	1907	−17.1	−23.5	2482	−17.3	−23.2				4410	−18	−23.2
9/10/2011	840	−14.8	−24.3	1246	−16.8	−24.7	1528	−17.2	−24.4				2656	−18.5	−24.5
10/6/2011	1478	−16.3	−23.3	3858	−18.1	−23.4	4033	−18.5	−23.8				5402	−18.5	−23.6
Date	Grassland														
	6 cm			12 cm			18 cm			24 cm					
	$p\text{CO}_2$	$\delta^{13}\text{C}$	$\delta^{13}\text{C}_r$	$p\text{CO}_2$	$\delta^{13}\text{C}$	$\delta^{13}\text{C}_r$	$p\text{CO}_2$	$\delta^{13}\text{C}$	$\delta^{13}\text{C}_r$	$p\text{CO}_2$	$\delta^{13}\text{C}$	$\delta^{13}\text{C}_r$	$p\text{CO}_2$	$\delta^{13}\text{C}$	$\delta^{13}\text{C}_r$
6/17/2010	1946	−11.9	−17.1	5403	−11.9	−16.5	6155	−12.8	−17.4	2961	−15.6	−21.0			
6/27/2011	2913	−13.4	−18.5	1544	−13.9	−20.0	5403	−14.3	−19.1	5952	−14.3	−19.0			
8/19/2011	580	−10.1	−17.5	730	−11.2	−18.5	765	−12	−19.8	791	−12.1	−19.7			
9/10/2011	619	−10.9	−19.0	807	−11.7	−18.9	890	−12.2	−19.3	1046	−12.5	−19.1			
10/6/2011	1823	−14.1	−19.9	2550	−14.6	−20.0	3038	−14.9	−20.1	3466	−15.3	−20.4			
Date	Streambed														
	11 cm			20 cm			23 cm			33 cm					
	$p\text{CO}_2$	$\delta^{13}\text{C}$	$\delta^{13}\text{C}_r$	$p\text{CO}_2$	$\delta^{13}\text{C}$	$\delta^{13}\text{C}_r$	$p\text{CO}_2$	$\delta^{13}\text{C}$	$\delta^{13}\text{C}_r$	$p\text{CO}_2$	$\delta^{13}\text{C}$	$\delta^{13}\text{C}_r$	$p\text{CO}_2$	$\delta^{13}\text{C}$	$\delta^{13}\text{C}_r$
6/17/2010	8464	−16.5	−21.2	7936	−15.4	−20.1	8325	−15.4	−20.0	13,696	−16.4	−20.9			
6/27/2011	4390	−14.8	−19.7	5752	−15.7	−20.5	7535	−16.3	−21.0	9034	−17	−21.7			
8/19/2011	977	−14.1	−22.0	1368	−15.1	−21.9	1795	−15.6	−21.8	2619	−16.2	−21.8			
9/10/2011	950	−14.2	−22.3	1484	−15.4	−22.1	2000	−15.9	−21.9	2708	−16.5	−22.1			
10/6/2011	2203	−15.6	−21.4	3377	−16.6	−21.9	4004	−16.9	−22.1	5455	−17.2	−22.2			
Date	Juniper stump														
	10 cm			15 cm			22 cm			29 cm					
	$p\text{CO}_2$	$\delta^{13}\text{C}$	$\delta^{13}\text{C}_r$	$p\text{CO}_2$	$\delta^{13}\text{C}$	$\delta^{13}\text{C}_r$	$p\text{CO}_2$	$\delta^{13}\text{C}$	$\delta^{13}\text{C}_r$	$p\text{CO}_2$	$\delta^{13}\text{C}$	$\delta^{13}\text{C}_r$	$p\text{CO}_2$	$\delta^{13}\text{C}$	$\delta^{13}\text{C}_r$
6/17/2010	1969	−17.3	−23.7	5443	−19.6	−24.7	15,120	−19.8	−24.4	20,090	−19.8	−24.3			
6/27/2011	2328	−17.7	−23.8	3106	−18.5	−24.2	4451	−18.9	−24.2	5330	−18.8	−23.9			
8/19/2011	539	−12.8	−27.4	760	−15.4	−26.6	1060	−16.7	−25.6	1375	−17.3	−25.0			
9/10/2011	576	−12.3	−24.0	728	−15	−26.4	985	−16.4	−25.6	1261	−17.2	−25.2			
10/6/2011	952	−15.7	−24.8	1218	−16.9	−25.0	1613	−17.6	−24.7	2022	−18	−24.5			

by grassland. The fraction of tree canopy coverage provides context for the carbon isotope measurements.

The source of carbon in soil CO₂ is better studied by considering $\delta^{13}\text{C}_r$ than $\delta^{13}\text{C}_s$ values (Cerling et al., 1991; Davidson, 1995). We suggest that if soils are the source of cave-air CO₂, then the advantage of studying $\delta^{13}\text{C}_r$ variations can and should be applied to cave-air CO₂ as well. Eq. (1) is appropriate for caves if CO₂ in caves or in the source region for cave-air CO₂ (i.e. soil or epikarst) reaches a steady-state exchange with the troposphere by diffusion. Calculation of $\delta^{13}\text{C}_r$ values allows comparison of the non-tropospheric components of cave-air CO₂ and soil CO₂ even if samples of these gases have different CO₂ concentrations. For instance, there is a consistent similarity between $\delta^{13}\text{C}_r$ values of cave-air CO₂ and tree-dominated soil CO₂ at COB (Table 8, Fig. 3B) but this similarity is not apparent if only the $\delta^{13}\text{C}$ values of CO₂ are compared (Table 8).

The consistent similarity between tree-dominated soil and cave-air $\delta^{13}\text{C}_r$ values and the consistent difference between grassland and cave-air $\delta^{13}\text{C}_r$ values suggests that the trees rather than grasses are the source of CO₂ in

COB. A similar comparison considering two endmember mixing (grassland and tree-dominated soil) at NB also suggests a dominantly (>80%) tree source for cave-air CO₂ (Fig. 3A). Tree canopies cover 15% and 60% of the surface above COB and NB, respectively. Therefore, the influence of trees on cave-air CO₂ is disproportionately large in comparison with their surface coverage. Furthermore, the $\delta^{13}\text{C}$ value of cave-air CO₂ is not controlled by the percentages of C₃ and C₄ biomass on the surface above these caves. We suggest that trees (or decomposition of OM from trees) control cave-air CO₂ because trees root deeper than grasses in these ecosystems (e.g. Jackson et al., 1999). In some ecosystems (e.g. prairie) C₄ grasses are deeper rooted than C₃ forbs. Therefore the conclusions from our study of juniper and oak savannah that grasses have little influence on cave CO₂ $\delta^{13}\text{C}$ values may not apply in ecosystems without trees and should be tested in ecosystems in which trees are not as deeply rooted as they are in our study areas.

We numerically simulated CO₂ production and transport by diffusion in soil and epikarst to investigate whether theory supports our empirically-derived conclusion that

Table 6
CO₂ concentrations and $\delta^{13}\text{C}$ values in WP cave-air.

Date	Outside air		1			2			3			4		
	pCO ₂	δ ¹³ C	pCO ₂	δ ¹³ C	δ ¹³ C _r	pCO ₂	δ ¹³ C	δ ¹³ C _r	pCO ₂	δ ¹³ C	δ ¹³ C _r	pCO ₂	δ ¹³ C	δ ¹³ C _r
10/23/2008	680	−13.2				5586	−21.0	−26.4				14,420	−16.9	−21.4
2/26/2009	314	−8.3				1397	−16.4	−23.0				2223	−17.4	−23.2
4/15/2009	370	−9.5				2220	−17.3	−23.2				3410	−19.4	−24.9
8/11/2009	393	−8.3	2151	−14.9	−20.7				2725	−15.9	−21.5			
8/1/2010						4841	−15.9	−20.8	5180	−16.4	−21.3			

Date	5			6			7			8		
	pCO ₂	δ ¹³ C	δ ¹³ C _r	pCO ₂	δ ¹³ C	δ ¹³ C _r	pCO ₂	δ ¹³ C	δ ¹³ C _r	pCO ₂	δ ¹³ C	δ ¹³ C _r
10/23/2008				13,860	−18.9	−23.5						
2/26/2009							5062	−18.0	−22.9			
4/15/2009										4020	−19.2	−24.5
8/11/2009	3451	−16.6	−22.0	4714	−16.8	−21.9						
8/1/2010	5017	−16.2	−21.1									

Date	10			11			12			13		
	pCO ₂	δ ¹³ C	δ ¹³ C _r	pCO ₂	δ ¹³ C	δ ¹³ C _r	pCO ₂	δ ¹³ C	δ ¹³ C _r	pCO ₂	δ ¹³ C	δ ¹³ C _r
10/23/2008	12,700	−19.0	−23.6	13,190	−19.0	−23.6	17,350	−19.4	−23.9	17,790	−18.4	−22.9
2/26/2009				4834	−18.1	−23.1	8174	−18.5	−23.2	9920	−18.7	−23.3
4/15/2009				4900	−19.4	−24.5				5500	−18.8	−23.8
8/11/2009				7812	−16.8	−21.6				12,300	−16.4	−21.0
8/1/2010	7347	−16.9	−21.6							15,592	−18.1	−22.6

Table 7
CO₂ concentrations and $\delta^{13}\text{C}$ values in WP soils.

Date	Grassland											
	10 cm			15 cm			20 cm			25 cm		
	$p\text{CO}_2$	$\delta^{13}\text{C}$	$\delta^{13}\text{C}_r$	$p\text{CO}_2$	$\delta^{13}\text{C}$	$\delta^{13}\text{C}_r$	$p\text{CO}_2$	$\delta^{13}\text{C}$	$\delta^{13}\text{C}_r$	$p\text{CO}_2$	$\delta^{13}\text{C}$	$\delta^{13}\text{C}_r$
8/1/2010	5735	−14.6	−19.3	8461	−15.1	−19.7	10,436	−15.4	−19.9	19,719	−15.6	−20.0
Date	Woodland											
	10 cm			15 cm			20 cm			25 cm		
	$p\text{CO}_2$	$\delta^{13}\text{C}$	$\delta^{13}\text{C}_r$	$p\text{CO}_2$	$\delta^{13}\text{C}$	$\delta^{13}\text{C}_r$	$p\text{CO}_2$	$\delta^{13}\text{C}$	$\delta^{13}\text{C}_r$	$p\text{CO}_2$	$\delta^{13}\text{C}$	$\delta^{13}\text{C}_r$
8/1/2010	6773	−18.1	−23.0	9461	−18.8	−23.6	8080	−18.6	−23.5	11,946	−19.4	−24.1

deeply rooted plants control cave-air CO₂ and therefore whether this conclusion might be applicable to ecosystems other than the ones studied here. We used the Forward-Time Central-Space (FTCS) method to solve Fick's second law of diffusion with a production term. Concentrations of CO₂ isotopologues were fixed at tropospheric values at the top of the soil and a zero gradient was established as the lower boundary condition. The effective diffusion coefficient for CO₂ was calculated as a function of free air porosity and pore space tortuosity (Bird et al., 1960; Quade et al., 1989). The diffusion coefficients for CO₂ isotopologues were calculated according to Craig (1954) and Jost (1960), following Cerling (1984). Errors associated with numerical approximation are insignificant at time steps ≤ 6 s. Simulated CO₂ concentrations and $\delta^{13}\text{C}_s$ values were used to calculate apparent $\delta^{13}\text{C}_r$ values, which were then compared with empirical $\delta^{13}\text{C}_r$ values. We simulated a steady-state, 10 m deep CO₂ profile through soil and epikarst assuming that CO₂ with a $\delta^{13}\text{C}$ value of -19‰ is respired in the

top 50 cm (typical soil depth on the Edwards Plateau is 20 cm) and CO₂ with a $\delta^{13}\text{C}$ value of -24‰ is respired at all depths below 50 cm (i.e. epikarst). Respiration rates were assumed to decrease exponentially with depth as shown in Fig. 5D such that 88% of the CO₂ respired in the soil–epikarst column came from the top 50 cm and the weighted mean $\delta^{13}\text{C}$ value of respired CO₂ was -19.6‰ . In the simulation, less than 3% of the CO₂ respired came from below 4 m. The free air porosity was set at 0.5 from 0 to 0.5 m, at 0.3 from 0.5 to 4 m and at 0.1 below 4 m. This simple scenario was intended to test the relative influence on cave-air CO₂ of shallow (grassland soil) versus deep (trees roots in epikarst) respiration in areas where tree roots extend at depth beneath grasslands. Calculated apparent $\delta^{13}\text{C}_r$ values approach -24‰ (the prescribed tree root value and the mean observed cave-air value) at 10 meters, which is the approximate depth of the shallowest cave (WP) we studied. The simulated, steady-state, vertical gradient of apparent $\delta^{13}\text{C}_r$ values (Fig. 5B, dotted line)

Table 8
CO₂ concentrations and $\delta^{13}\text{C}$ values in COB cave-air and soils.

Date	Popcorn room			North room streambed woodland (61 cm)			Streambed woodland (61 cm)			Grassland 1 (55 cm)			Grassland 2 (70 cm)		
	$p\text{CO}_2$	$\delta^{13}\text{C}$	$\delta^{13}\text{C}_r$	$p\text{CO}_2$	$\delta^{13}\text{C}$	$\delta^{13}\text{C}_r$	$p\text{CO}_2$	$\delta^{13}\text{C}$	$\delta^{13}\text{C}_r$	$p\text{CO}_2$	$\delta^{13}\text{C}$	$\delta^{13}\text{C}_r$	$p\text{CO}_2$	$\delta^{13}\text{C}$	$\delta^{13}\text{C}_r$
9/5/2008							14,432	−20.0	−24.6	28,012	−15.6	−20.1	24,313	−16.7	−21.1
11/5/2008	9200	−19.0	−23.7							8488	−14.3	−18.9	8004	−15.6	−20.2
1/8/2009	9463	−19.2	−23.9	8703	−19.2	−24.0	2417	−18.0	−24.0	4792	−16.6	−21.6	6683	−16.9	−21.7
2/7/2009	8107	−18.8	−23.6	8327	−18.8	−23.6	2191	−17.7	−23.9	4221	−16.0	−21.1	5436	−16.7	−21.6
3/6/2009	8535	−19.1	−23.9	8595	−19.2	−23.9	2296	−17.5	−23.6	4759	−15.4	−20.3	4987	−16.1	−21.0
4/3/2009	8484	−19.5	−24.3	8221	−19.6	−24.4	1830	−17.7	−24.4	3758	−15.3	−20.4	5071	−16.2	−21.1
5/3/2009	9657	−19.1	−23.8	7439	−19.3	−24.2	1795	−17.0	−23.6	3761	−14.1	−19.1			
6/6/2009	9776	−19.6	−24.4	10,265	−19.5	−24.2	2542	−17.7	−23.6	4938	−14.0	−18.7	6099	−14.9	−19.6
8/1/2009							10,310	−19.3	−24.0						
1/16/2010	8238	−19.5	−24.3	8017	−19.6	−24.5	1462	−17.1	−24.4	2151	−16.1	−22.0	2323	−16.3	−22.1
4/24/2010	8421	−19.5	−24.3	8421	−19.9	−24.7	5178	−19.8	−25.0	8140	−16.4	−21.1	7048	−16.4	−21.2
5/22/2010	7892	−18.5	−23.3	7328	−18.3	−23.1	5065	−18.1	−23.1	8086	−13.9	−18.4	9760	−15.2	−19.8
7/28/2010	11,281	−18.8	−23.5	11,609	−18.9	−23.5	7436	−18.1	−22.9	11,128	−14.8	−19.3	12,403	−15.5	−20.0

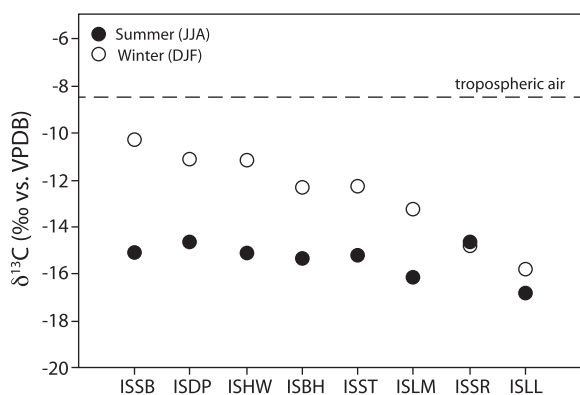


Fig. 2. Mean measured $\delta^{13}\text{C}$ values of cave-air CO₂ along transects into IS during our 18 month study. During the winter, the $\delta^{13}\text{C}$ values of cave-air CO₂ decrease from near tropospheric values at the cave entrance to -16‰ at the site furthest from the entrance. The gradient is shallower during the summer. These patterns are consistent with ventilation of IS by cold, dense tropospheric air during the winter. Locations of cave-air CO₂ sampling are shown in Fig. 1. IS refers to Inner Space in all sample names, the last two letters of the sampling are abbreviations as follows: SB = Soda Straw Balcony, DP = Drapery, HW = Hallway, BH = Borehole, ST = Flowing Stone of Time, LM = Lake of the Moon, SR = Squid Room, LL = Lunar Landscape.

demonstrates that respiration in the epikarst, rather than the surface soil, may control the $\delta^{13}\text{C}_r$ value of cave-air CO₂. Even low respiration in the epikarst can have a significant influence on cave-air CO₂ $\delta^{13}\text{C}$ values, especially if the effective diffusion coefficient for CO₂ in the deep epikarst is small. This conclusion is consistent with the speleogenesis model conceived by Wood (1985) in which low rates of deep vadose zone respiration (as opposed to soil respiration) are responsible for the high CO₂ concentrations in the deep vadose zone. Consideration of advection in numerical gas transport simulations would improve the application to ventilated caves.

Profiles of the fmc in CO₂ were also simulated using the FTCS method. Small fluxes of CO₂ from oxidation of passive soil carbon (hundreds to thousands of years old, e.g. O'Brien and Stout, 1978) deep in the epikarst might explain the relatively low radiocarbon specific activity in COB cave-air CO₂. The importance of old organic matter oxidation on speleothem carbon isotope compositions has been recently documented (Oster et al., 2010; Rudzka et al., 2011). One scenario that satisfies the observations made in this study is oxidation of ~ 1000 A.D. organic matter below 4 m, at a rate small enough that the weighted mean fmc of CO₂ respired in the entire soil and epikarst column is 1.075 (i.e. nearly equivalent to the observed value in the soil). The wetter-than-modern climate during the medieval warm period (e.g. Davis, 1994; Leavitt, 1994) might explain the presence of old organic matter at depth in the epikarst with $\delta^{13}\text{C}$ values similar-to-modern despite the higher-than-modern $\delta^{13}\text{C}$ values of preindustrial tropospheric CO₂. An old organic carbon source for cave-air CO₂ is consistent with the slow change in $\delta^{13}\text{C}$ values of speleothem calcite observed in the Cold Water Cave speleothem record (Dorale et al., 1992). Fractions of carbon from limestone in COB cave-air CO₂ that are larger than a couple of percent are difficult to explain given the close agreement between soil and cave-air CO₂ $\delta^{13}\text{C}_r$ values, unless soil CO₂ also contains limestone carbon. It is also possible that cave-air is equilibrated with a large volume of old groundwater (volume of air filled cave $\ll$ volume of groundwater). In this case, preferential recharge through woodland dominated streambeds followed by aging of carbon in the saturated zone could explain the observations. Measurements of the radiocarbon activity of drip water and the age of groundwater in this region would help distinguish between these two conceptual models. If the isotopic composition of CO₂ in COB air is controlled by groundwater, then the deeply rooted plants or the plants that grow in the streambeds must control the $\delta^{13}\text{C}$ value of groundwater dissolved inorganic carbon (DIC). This idea should be tested in other aquifers.

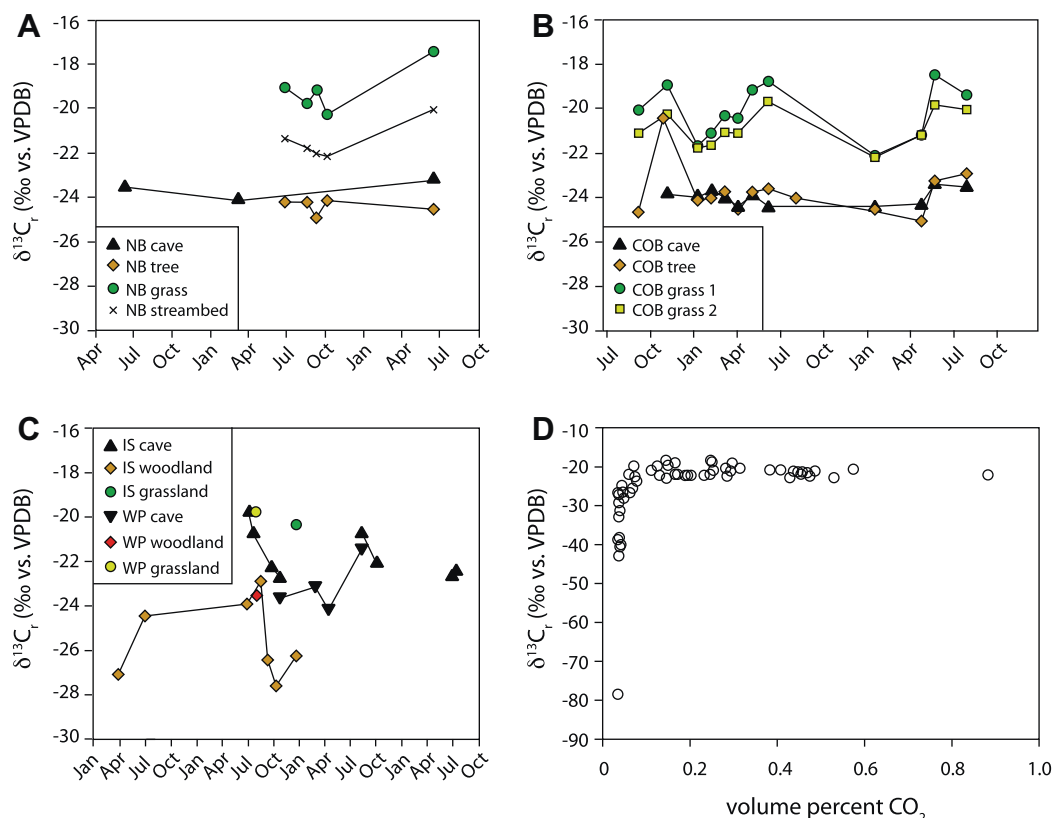


Fig. 3. Time series of calculated soil and cave-air CO_2 $\delta^{13}\text{C}_t$ values. Soils dominated by trees and by grasses are indicated by diamonds and by circles, respectively. (A) Natural Bride Caverns (NB). X's represent CO_2 in a gravelly streambed above the cave. (B) Cave of the Bells (COB). (C) Inner Space Caverns (IS) and Whirlpool Cave (WP). (D) Relationship between CO_2 concentrations and calculated $\delta^{13}\text{C}_t$ values for all sites in IS. Low $\delta^{13}\text{C}_t$ values that occur when cave-air CO_2 is below 1500 ppmV likely result from kinetic isotope fractionation during degassing of CO_2 -supersaturated drip waters. $\delta^{13}\text{C}_t$ values for times when IS CO_2 was below 1500 ppmV (primarily winter) are therefore not included in the comparison with soil CO_2 in panel C. These results indicate that CO_2 in NB and COB comes primarily from tree-dominated soils whereas the CO_2 in IS and WP is primarily from grasslands during the summer and from both grasslands and woodlands during the winter. The uncertainty associated with the $\delta^{13}\text{C}_t$ values plotted is $\pm 0.2\text{‰}$.

If respiration of deep and possibly old organic carbon controls the carbon isotope composition of cave-air CO_2 , then it probably also influences the carbon isotope composition of seepage waters and speleothem calcite. If true, then speleothems may provide records of the carbon isotope composition of deeply rooted plants and/or the presence/absence of deeply rooted plants. In addition, radiocarbon calibrations should take into account that carbon may have aged in the soil or epikarst for a 1000 years or longer before being incorporated into speleothem calcite, even in the absence of incorporation of limestone carbon. Measurements of the carbon isotope composition of cave waters and harvested cave calcite are required to test these ideas.

Calculated $\delta^{13}\text{C}_t$ values of cave-air CO_2 in IS during the winter (-25 to -80‰ , Table 2) are too low to be interpreted as soil-derived. Instead, the low $\delta^{13}\text{C}_t$ values that occur when cave-air CO_2 concentrations are below 1500 ppmV CO_2 (Fig. 3D) are likely controlled by preferential degassing of $^{12}\text{CO}_2$ from CO_2 -supersaturated drip water (Spötl et al., 2005). Therefore, water likely transports a substantial fraction of CO_2 into IS during the winter. Unfortunately, this process obscures the determination of carbon

sources using winter $\delta^{13}\text{C}_t$ values of cave-air CO_2 in IS. The sources of cave-air CO_2 in IS can, however, be assessed at other times of year.

During the summer, $\delta^{13}\text{C}_t$ values of cave-air CO_2 in IS and WP are similar to $\delta^{13}\text{C}_t$ values of grassland soil CO_2 measured at WP and NB (we did not measure $\delta^{13}\text{C}_t$ values of grassland soil CO_2 at IS during the summer). We therefore suggest that grasslands are the primary source for cave-air CO_2 during the summer at IS and WP, which is consistent with the large area of contiguous grassland above these caves. In comparison to IS, the savannah ecosystem above COB has a similar percentage of tree cover ($\sim 15\%$), but the trees are more evenly distributed at COB whereas at IS the trees mostly occur in woodlands located over distal corners of the cave (Fig. 1). The trees are more clustered above WP than COB and a large patch of grassland occurs above the region within WP where most of the cave-air CO_2 samples were collected (Fig. 1). Our one-dimensional numerical simulation indicates that deeply rooted trees will dominate the contribution to cave-air CO_2 if they grow close enough together. However, larger distances between trees, like those at IS and WP, should increase the influence of shallower

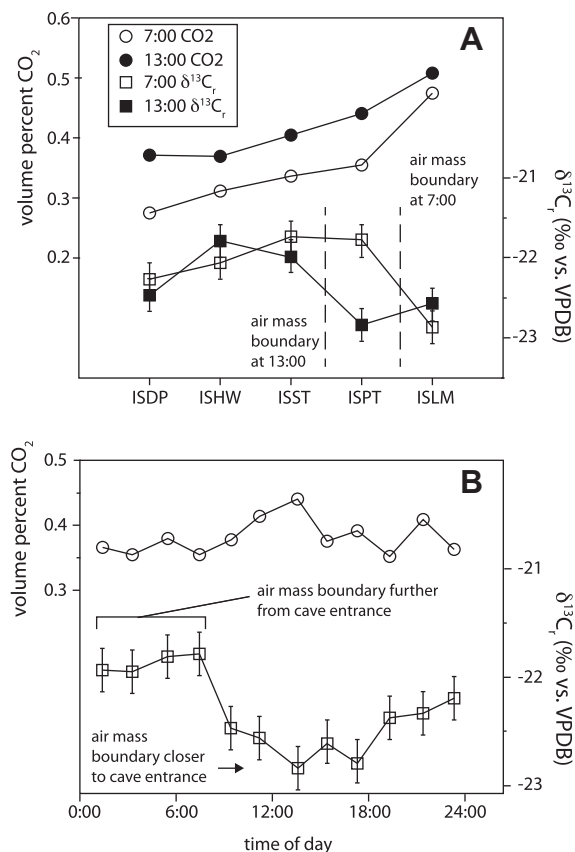


Fig. 4. CO₂ concentrations and δ¹³C_r values of cave-air CO₂ in IS during the July 2010 24-h study. (A) Comparison of transects into IS during the morning and early afternoon. (B) Diurnal variations in the concentration and δ¹³C_r value of cave-air CO₂ at sampling site ISPT. Air masses (boundaries defined by dotted lines in A) with distinct δ¹³C_r values move back and forth across ISPT with a period of approximately 24 h. This process is not recorded by cave-air CO₂ concentrations alone. ISPT = Inner Space Painting.

grassland soil respiration on the δ¹³C value of cave-air CO₂. A two-dimensional CO₂ production and transport model would help test this idea.

A decrease of ~3‰ in the δ¹³C_r value of cave-air CO₂ occurs from July through October in IS. This decrease is not related to kinetic fractionation during degassing, as demonstrated by the poor correlation between δ¹³C_r values and cave-air CO₂ concentrations during this time period ($r^2 = 0.02$, $p = 0.77$, $n = 8$). The decrease in δ¹³C_r values may instead result from an increase in the relative contribution of woodland soil CO₂ to the cave, a decrease in the δ¹³C value of CO₂ respired in all soils above the cave, or perhaps both. The difference between δ¹³C_r values of cave-air CO₂ in WP and IS on the same days in October and August were 0.9‰ and 0.8‰, respectively (Fig. 3C). The δ¹³C_r values of cave-air CO₂ in WP may therefore be used to approximate the source of CO₂ in IS during the winter. Comparison of winter δ¹³C_r values of cave-air CO₂ in WP (+1‰ to account for the difference observed in October and August) with winter δ¹³C_r values of grassland and oak canopy soil CO₂ above IS suggests there is

a 40% contribution of CO₂ from trees. Although this is a smaller fraction than the contribution from trees at COB and NB, it is large in comparison with the fraction of the surface covered by tree canopies. Further monitoring is required to better quantify the contribution of trees to the CO₂ in IS.

Human breath is another potential source of CO₂ in caves. The average δ¹³C value of CO₂ in breath of North Americans is between −21‰ and −20‰ (Schoeller et al., 1980), which is intermediate between δ¹³C_r values of grassland and tree-dominated soil CO₂ and is similar to δ¹³C_r values of mean cave-air CO₂ in IS during the summer-fall transition. The δ¹³C values of CO₂ in human breath measured in this study are similar to the values reported by Schoeller et al. (1980) and are significantly higher than the δ¹³C_r value of cave-air CO₂ in COB, NB and WP ($p < 0.002$ in each case). Furthermore, δ¹³C_r values of cave-air CO₂ observed at NB (a tourist cave) are similar to values at COB and WP (wild caves with relatively few visitors). Therefore human breath is probably not responsible for the tree-biased δ¹³C_r values of cave-air CO₂ in the caves we studied. High-resolution temporal variations in cave-air CO₂ concentrations in IS do not peak during the passage of tour groups (Cowan, 2010), further supporting the conclusion that human breath is of minor importance for CO₂ in these caves.

6.3. CO₂ transport into caves

The mechanism of CO₂ transport into caves has not been widely studied. The similarities between δ¹³C_r values of soil CO₂ and cave-air CO₂ observed in this study suggest that the majority of CO₂ is transported into these caves as a gas and not dissolved in water. If carbon isotope equilibrium is maintained between CO₂ and DIC during degassing from the water, the δ¹³C value of the first CO₂ emitted into the cave atmosphere would equal δ¹³C_s not the δ¹³C value of CO₂ respired in the soil (δ¹³C_{r-soil}; because the water equilibrated in the epikarst with pore space CO₂, not respired CO₂). The δ¹³C value of degassed CO₂ would increase with the fraction of total carbon degassed because $\alpha_{\text{CO}_2-\text{HCO}_3^-} < 1$; 100% degassing would result in cave-air CO₂ δ¹³C_r values equal to the δ¹³C value of DIC in equilibrium with soil CO₂. The occurrence of kinetic isotope effects during degassing would result in δ¹³C_r values of cave-air CO₂ that are lower than δ¹³C_r values of soil CO₂, as observed in IS during the winter. Therefore, the observed similarity between tree-dominated soil and cave δ¹³C_r values requires either (1) equilibrium degassing of a small fraction of cave water DIC and advective mixing of cave-air to prevent carbon isotope fractionation by diffusion of CO₂ or (2) transport of CO₂ into caves as a gas. Under conditions required for (1), all speleothem calcite might precipitate in isotope equilibrium with soil CO₂ and vadose water, which would be ideal for paleoclimate reconstruction. We, however, think (2) is more likely, given evidence presented here and previously for kinetic isotope fraction during degassing of CO₂ from cave drip waters (e.g. Spötl et al., 2005). Precise measurement of oxygen concentrations in cave-air would help distinguish between CO₂ transport into caves in water or as a gas (Halbert, 1982). Depletions of O₂ in cave-air

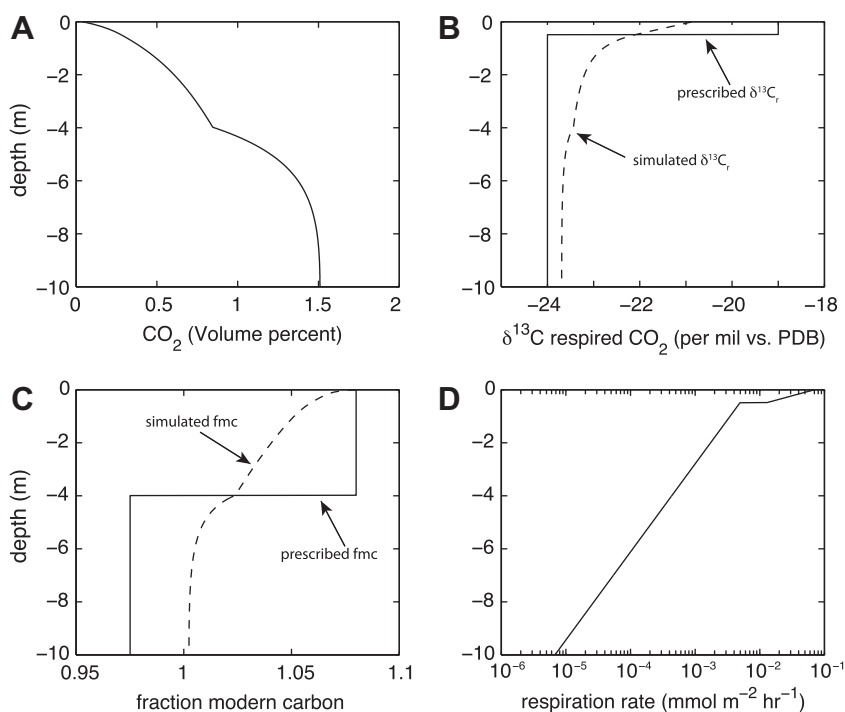


Fig. 5. Simulations of pore space CO₂ in a 10 m column of soil and epikarst. A numerical production-diffusion model was used to simulate steady-state profiles of CO₂. (A) CO₂ concentrations, (B) $\delta^{13}\text{C}$ values of respired CO₂. The solid line shows values prescribed as model input; the dashed line shows simulated $\delta^{13}\text{C}_r$ values calculated from model output using equation 1 from the text. The simulated $\delta^{13}\text{C}_r$ values can be compared with $\delta^{13}\text{C}_r$ values calculated from measured concentrations and $\delta^{13}\text{C}$ values of soil and cave-air CO₂. (C) The fraction modern carbon (fmc) in pore space CO₂. The solid line shows values prescribed as model input; the dashed line shows values calculated from model output. (D) Respiration rates used as model input. Free air porosity was prescribed as follows: 0–0.5 m, porosity = 0.5; 0.5–2 m, porosity = 0.3; 2–10 m, porosity = 0.1. The tortuosity factor and temperature were held constant at 0.6 and 15 °C, respectively. CO₂ respired below 4 m was assigned a fmc of 0.87, equivalent to the fmc in 2009 of organic carbon removed from the atmosphere in 1000 A.D.

below tropospheric O₂ concentrations that balance the enrichment of cave-air CO₂ above tropospheric CO₂ concentrations would indicate gas phase transport between soil and cave whereas elevated cave-air CO₂ concentrations without compensatory O₂ consumption would indicate that CO₂ enters caves by degassing from water.

6.4. Seasonal variation of cave-air CO₂

The concentration of CO₂ in the cave atmosphere at a given point in time is the result of a balance between CO₂ fluxes into the cave from external sources and CO₂ fluxes out by removal mechanisms. Seasonal variations in cave-air CO₂ are the result of fluctuations in the magnitude of these fluxes. However, it is difficult to conclusively determine the dominant cause of cave-air CO₂ change because the source and sink strengths both change seasonally. $\delta^{13}\text{C}$ values alone are insufficient to distinguish between a reduced soil respiration flux or increased ventilation because both processes would result in an increased tropospheric CO₂ to soil-respired CO₂ ratio and therefore higher cave-air CO₂ $\delta^{13}\text{C}$ values.

Wong and Banner (2010) investigated the effects of vegetation removal over cave NBS. Their work provides an indirect means of resolving whether changes in soil respiration rate or changes in ventilation strength have a larger

effect on cave-air CO₂. Cave-air CO₂ was measured over a period of 5 years: 2 years with natural surface vegetation followed by 3 years over which a large portion of vegetation cover, primarily Juniper trees, above a section of the cave was removed (Wong and Banner, 2010). Within NBS, they found immediate and significant decreases in cave-air CO₂ concentrations and dampened summer CO₂ concentrations in the years following vegetation removal. The large effects of vegetation removal on cave-air CO₂ suggest that root respiration, specifically juniper root systems, are a significant source of CO₂ to the cave (Wong and Banner, 2010). This is consistent with our interpretations based on carbon isotope ratios of cave and soil CO₂. The lack of any observable response of cave-air CO₂ to brush removal during the winter months suggests that winter CO₂ decreases are primarily controlled by CO₂ removal through ventilation rather than decreases in soil respiration rate. It follows that soil/epikarst respiration controls how high CO₂ gets during the summer months and that density-driven ventilation controls how low CO₂ gets during the winter months. Lower summer cave-air CO₂ concentrations in IS than in the other caves may be the result of the shallow depths of respiration in the grassland-dominated soils above the cave or to the occasional use of ventilation shafts.

If all other factors are held equal, then shallower depths of respiration results in lower pore space CO₂

concentrations. Therefore, when compared with other caves we studied, shallow depths of respiration in the contiguous grasslands above IS may explain the lower summer maximum CO₂ concentrations in this cave. Furthermore, the presence of contiguous grassland and the associated shallow respiration might limit speleothem growth rates because epikarst CO₂ concentrations control the maximum Ca²⁺ concentrations of water entering caves. Likewise, temporal variation in the rooting depth of vegetation might influence the growth rate of speleothems through time.

6.5. Diurnal cave-air CO₂ cycle

Lateral transects into IS suggest that distinct air masses identifiable using measured $\delta^{13}\text{C}$ values and CO₂ concentrations exist within the cave (Fig. 4). The air mass furthest from the entrance of the cave, and closest to the woodland vegetation above the cave, has a $\delta^{13}\text{C}_r$ value of approximately -23‰ whereas an air mass in the vicinity of ISST has a $\delta^{13}\text{C}_r$ value about 1‰ higher (Fig. 4). The air mass closest to the entrance has an intermediate $\delta^{13}\text{C}_r$ value. The transitions between these air masses are relatively abrupt (~ 75 m) and their positions within the cave oscillate with a period of approximately 24 h; the transitions were furthest from the entrance at 7:00 and closest to the entrance at 13:00 (CST) during our 24 h monitoring study. Movement of the transitions between the air masses across sites ISPT and ISHW results in diurnal variation in $\delta^{13}\text{C}_r$ values at individual locations, which are especially pronounced at ISPT (Fig. 4). This diurnal movement of the air in and out of the cave is likely controlled by atmospheric tides (Volland, 1997) as concluded by Cowan (2010) for central Texas caves, whereby decreases in atmospheric pressure pull air from deeper in a cave toward its entrance and vice versa. The differences in $\delta^{13}\text{C}_r$ values among the air masses in IS likely result from differences in vegetation above the cave. The vegetation at the entrance to the cave is dominated by forbs and oak trees (Table 3 and Fig. 1) and a patch of woodland occurs above site ISLM, whereas mid-cave-air mass with relatively high $\delta^{13}\text{C}_r$ values is under grassland and Interstate Highway 35.

7. CONCLUSIONS

- (1) CO₂ is transported as a gas into the caves studied, except in Inner Space Cavern during the winter when a significant amount of CO₂ degasses from cave water. Calculating $\delta^{13}\text{C}$ values of respired CO₂ is therefore useful for investigating the sources of CO₂ in cave-air.
- (2) Deeply rooted vegetation, if present, supplies the majority of the CO₂ to caves even if the rates of deep respiration are low and the fraction of surface cover by deeply rooted plants is small, especially if the epikarst has a low porosity.
- (3) The distribution of vegetation is more important than the fraction of coverage in determining the relative contribution of CO₂ from different types of plants to cave-air. Large contiguous areas of shallowly

rooted vegetation can overcome the bias against shallow respiration in the $\delta^{13}\text{C}$ value of deep pore space CO₂.

- (4) Oxidation of old organic matter in the epikarst or equilibration of cave-air with large volumes of old groundwater may result in radiocarbon activities of cave-air CO₂ that are lower than the activities in soil CO₂.
- (5) Variations in the $\delta^{13}\text{C}$ value of CO₂ in cave-air support the ideas that density-driven ventilation is a sink for cave-air CO₂ during the winter and that diurnal barometric pumping during the summer shifts air mass inside caves.

These considerations lay the groundwork for more accurate interpretations of speleothem-based paleoclimate proxies, particularly the carbon isotope composition of speleothem calcite.

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